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Amendments to the Claims

This listing of claims replaces all prior versions of claims in the application:

1. (Currently amended) A pharmaceutical dosage form suitable for oral administration comprising a molded microcellular polymeric material and a pharmaceutically acceptable active agent, and

wherein the molded microcellular polymeric material is a non-thermosetting polymerized plastics material comprised of at least one polyol selected from lactitol, xylitol, erythritol, sorbitol, maltitol, or mannitol, or combinations thereof; and

at least one of

a) a non-thermosetting modifier selected from a starch, maltodextrin, a dextrose equivalent, polyalditol, a hydrogenated starch hydrosylate, or a mixture thereof; and/or

b) a non-thermosetting polymer selected from carboxymethyl cellulose sodium, methyl cellulose, ethylcellulose, hydroxyethylcellulose (HEC), hydroxypropylmethyl cellulose (HPMC), hydroxypropylmethyl cellulose phthalate, cellulose acetate phthalate, noncrystalline cellulose, starch and its derivatives, and sodium starch glycolate or mixtures thereof; and

c) optionally a sweetener, a disintegrant, a binder, a lubricant or an opacifier; and

wherein the molded microcellular polymeric material and pharmaceutically active agent are injection molded into the pharmaceutical dosage form.

2. -3. (Cancelled)

4. (Previously presented) The pharmaceutical dosage form according to claim 1 wherein the non-thermosetting polymerized plastics material contains at least one polyol, and at least one non-thermosetting modifier.

5. – 6. (Cancelled)

7. (Previously presented) The pharmaceutical dosage form according to claim 1 wherein the starch is pregelatinized corn starch, corn starch, potato starch, rice starch,